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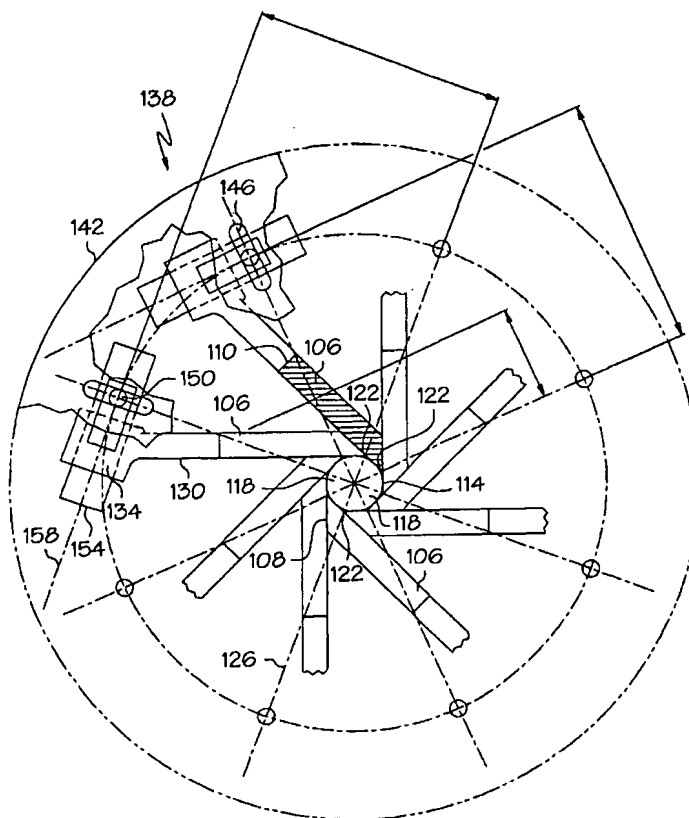
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(54) Title: A METHOD FOR CONTRACTING, LOADING OR CRIMPING SELF-EXPANDING AND BALLOON EXPANDABLE STENT DEVICES



(57) Abstract: An apparatus for manipulating a medical device is formed of at least three coupled movable blades which are disposed about a reference circle to form an aperture whose size may be varied. The aperture capable of being sized to contain a medical device. Each blade is in communication with an actuation device which is capable of moving the blade to alter the size of the aperture. Each blade includes a single radial point which a) lies on the circumference of the reference circle prior to movement of the blade, and b) may be moved only along a radius of the reference circle on movement of the blade.

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A METHOD FOR CONTRACTING, LOADING OR CRIMPING SELF-EXPANDING AND BALLOON EXPANDABLE STENT DEVICES

BACKGROUND OF THE INVENTION

5 This invention relates to an apparatus and a method for reducing in size a medical device such as a stent, stent-graft, graft, or vena cava filter. The apparatus may be used in particular for fastening a medical device onto a catheter.

 Medical devices such as stents, stent-grafts, grafts, or vena cava filters and catheters for their delivery are utilized in a number of medical procedures and
10 situations, and as such their structure and function are well known.

 A stent, for example, is a generally cylindrical prosthesis introduced via a catheter into a lumen of a body vessel in a configuration having a generally reduced diameter and then expanded to the diameter of the vessel. In its expanded configuration, the stent supports and reinforces the vessel walls while maintaining the vessel in an
15 open, unobstructed condition.

 Stents are typically inflation expandable or self-expanding. Self expanding stents which are constrained by a sheath or other restraining means, must be provided in a reduced diameter.

 An example of a stent described in PCT Application No. 960 3092 A1,
20 published 8 February 1996.

 In advancing a stent through a body vessel to the deployment site, the stent must be able to securely maintain its axial position on the delivery catheter, without translocating proximally or distally, and especially without becoming separated from the catheter. Stents that are not properly secured or retained to the catheter may
25 slip and either be lost or be deployed in the wrong location. The stent must be crimped in such a way as to minimize or prevent altogether distortion of the stent and to thereby prevent abrasion and/or reduce trauma of the vessel walls.

 In the past, this crimping or size reduction has been done by hand often resulting in the application of undesired uneven forces to the stent. Such a stent must
30 either be discarded or re-crimped. Stents which have been crimped or otherwise reduced in size multiple times can suffer from fatigue and may be scored or otherwise marked which can cause thrombosis. A poorly crimped stent can also damage the underlying balloon.

Recently, stent crimping devices have been disclosed in US 5,546,646 to Williams et al, US 5,183,085 to Timmermans et al., US 5,626,604 to Cottone, Jr., US 5,725,519, US 5,810,873 to Morales, WO 97/20593 and WO 98/19633.

A cam actuated stent crimper, shown in Figure 1, employs a plurality of
5 arc-shaped or curved slots with semi-circular ends, disposed such that each slot or cam engages a cam follower bearing 22. The arc-shaped or curved surfaces of the slots are inclined to be non-concentric relative to the axis of rotation 26, and therefore rotation of the cam plate 28 transmits equal radial displacements to the cam follower bearings 22, to simultaneously actuate a like number of linear bearings 24, which have their
10 corresponding linear tracks or rails mounted on a fixed plate. As shown in Fig.1 the cam plate rotary drive 29 comprises a pneumatic cylinder mounted on a pivot or trunnion, arranged with the cylinder rod connected rotatably to a short arm fixed rigidly to the cam plate. Accordingly, linear motion produced by the pneumatic cylinder translates into controllable arcs of motion of the circular cam plate, which has a
15 projecting V-shaped profile on its outer edge in rolling engagement with three equally spaced rollers with mating inverse V-shaped profiles to provide precise rotatable support to the cam plate. Depending on the direction of rotation, the linear slides which each carry a radially disposed crimping blade, are either moved inwards to apply a crimping force to the stent, or outwards to release the stent. Also when crimping,
20 depending on the degree of rotation of the cam plate, a specific radial crimping displacement may be obtained to match the diametral reduction required for any particular stent.

All US patents and applications and all other published documents mentioned anywhere in this application are incorporated herein by reference in their
25 entirety.

BRIEF SUMMARY OF THE INVENTION

It would be desirable to produce a device capable of crimping a stent uniformly while minimizing the distortion of and scoring and marking of the stent due
30 to the crimping. The present invention is directed to that end.

The present invention is particularly concerned with the crimping and otherwise reducing in size of inflation expandable stents, self-expanding stents and other expandable medical devices. For the purpose of this disclosure, it is understood

that the term 'stent' includes stents, stent-grafts, grafts and vena cava filters. It is also understood that the term 'crimping' refers to a reduction in size or profile of a stent.

In the description that follows it is understood that the invention contemplates crimping a medical device either directly to a catheter tube or to a catheter
5 balloon which is disposed about a catheter tube. When reference is made to crimping a medical device to a catheter, a balloon may be situated between the medical device and the catheter tube or the medical device may be crimped to a region of a catheter tube directly. The invention also contemplates crimping a stent in the absence of a catheter to reduce the stent in size.

10 The present invention is directed, in one embodiment, to an apparatus for reducing a medical device in size. Desirably, the medical device is a stent, a stent-graft, a graft or a vena cava filter, whether self-expandable, balloon expandable or otherwise expandable, although the inventive apparatus may also be employed with any other suitable, generally tubular medical device which must be reduced in size.

15 The inventive apparatus comprises at least three coupled movable blades disposed about a reference circle to form an aperture whose size may be varied. Each blade is in communication with an actuation device which is capable of moving the blade to alter the size of the aperture. Each blade includes a single radial point on the surface of the blade which a) lies on the circumference of the reference circle prior to
20 movement of the blade, and b) may be moved only along a radius of the reference circle on movement of the blade.

The apparatus further includes an actuation device which comprises a cam and a plurality of linear slide devices. Each linear slide device is in communication with a blade. Each of the linear slide devices is also in mechanical communication with
25 the cam. Rotation of the cam results in linear translation of the slide device and blade, such that the slide device moves along an axis parallel to the radius on which the radial point of the blade lies or along the radius itself.

The invention is also directed to an apparatus similar to that described above, with blades disposed about a reference tube to form a tubular aperture whose size
30 may be varied. Each blade is in communication with an actuation device which is capable of moving the blade to alter the size of the tubular aperture. Each blade includes a single line which a) lies on the surface of the reference tube prior to

movement of the blade, and b) may be moved only along a radial plane of the reference tube on movement of the blade.

The inventive apparatus finds particular utility in crimping a medical device such as those mentioned above to a catheter or to a balloon disposed about a catheter.

The inventive apparatus also finds utility in reducing the diameter of a medical device such as those mentioned above prior to crimping.

The invention is also directed to a method of manipulating a medical device which comprises the steps of providing the medical device and providing at least three blades capable of applying a radial inward force. The blades are disposed about a reference circle to form a shrinkable aperture. A medical device such as a stent is placed into the shrinkable aperture and the blades simultaneously moved inward to apply a radial inward force to the medical device. The blades are constructed and arranged such that each blade has a single point which a) lies on the circumference of the reference circle prior to movement of the blade, and b) is moved along a radius of the reference circle on movement of the blade.

The inventive apparatus may also be used as a variable size balloon mold. To that end, the invention is further directed to a method of molding a medical balloon. In the practice of the method, a balloon preform prepared through any suitable technique known in the art is provided. The preform is placed in an apparatus which has a shrinkable tubular aperture formed by at least three movable blades disposed about a reference tube. The blades are constructed and arranged such that each blade has a single line which a) lies on the surface of the reference tube prior to movement of the blade, and b) is moved along a radial plane of the reference tube on movement of the blade. The aperture may be set to a predetermined size prior to placement of the preform therein or after placement of the preform therein. An inflation fluid is supplied to the balloon preform to expand the balloon preform until it contacts the blades. The preform may optionally be heated prior to, during or after the blowing step. The thus formed balloon is then pressure relieved and removed from the apparatus.

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWING(S)

Figure 1 shows a perspective view of a stent crimper;

Figure 2a is a schematic front view of an embodiment of the inventive apparatus;

Figure 2b is a schematic front view of the embodiment of Fig. 2a after the stent has been reduced in size;

5 Figures 3a and 3b are schematics of blades;

Figure 3c is a partial schematic front view of an embodiment of the inventive apparatus employing the curved blades of Figure 3b;

Figure 4a is a partial front view of an embodiment of the inventive apparatus;

10 Figure 4b is a partial front view of an embodiment of the inventive apparatus;

Figure 4c shows a side view of the embodiment of Figure 4b taken along lines 4c-4c;

Figure 5a shows a partial front view of another embodiment of the
15 inventive apparatus;

Figure 5b shows a link connected to a blade;

Figure 6 is a schematic, perspective view of an embodiment of the inventive apparatus;

Figure 7 shows a partial view of the embodiment of Figure 6;

20 Figures 8a and 8b are partial side elevational views of an embodiment of the inventive apparatus taken along a radial plane during the size reduction process;

Figure 8c is a partial side elevational view of an embodiment of the inventive apparatus taken along a radial plane following crimping of a stent to a catheter;

25 Figure 9 is a diagrammatic side elevational view of an embodiment of the inventive apparatus;

Figure 10 is a partial side elevational view of an embodiment of the inventive apparatus taken along a radial plane of an embodiment of the invention consisting of three individual apparatuses arranged sequentially;

30 Figure 11 is a schematic showing a stent being reduced in size and loaded into a sheath;

Figure 12 is a partial side elevational view of an embodiment of the inventive apparatus taken along a radial plane showing a balloon that has been molded with the inventive device; and

Figure 13 is a partial side elevational view taken along a radial plane
5 showing a stepped balloon that has been molded with the inventive device.

DETAILED DESCRIPTION OF THE INVENTION

While this invention may be embodied in many different forms, there are described in detail herein specific preferred embodiments of the invention. This
10 description is an exemplification of the principles of the invention and is not intended to limit the invention to the particular embodiments illustrated.

As shown generally at 100 in Figures 2a and 2b, the inventive apparatus comprises eight coupled blades 106 disposed about a reference circle 114 to form an aperture 118 whose size may be varied. The apparatus may comprise as few as three
15 blades and as many as sixteen or more blades. Desirably, the apparatus will have four or more blades and more desirably, eight or more blades. The maximum number of blades is limited only by how many blades can physically be coupled together under the relevant size constraints. As the number of blades is increased, the profile of the aperture and hence of the medical device following reduction in size, becomes
20 smoother. Figure 2b shows the apparatus of Figure 2a after the stent has been reduced in size.

Blades 106 as shown in Figure 3a have an inner end 108 which is desirably beveled 111 so as to mesh with adjacent blades and an outer end 110 which is displaced from aperture 118. Aperture 118 is polygonal. Blades 106 may also be shaped
25 with a curved end 112, as shown in Figures 3b and 3c so as to form a substantially circular shaped aperture, when the aperture is fully closed.

Each blade 106 includes a single radial point 122 which lies on a radial line 126 of reference circle 114 prior to movement of blade 106 and which may be moved only along the radius 126 of reference circle 114 on movement of blade 106.
30 Desirably, the single radial point 122 will be disposed at the tip of the blade adjacent to beveled end 111.

In the embodiment shown in Figure 4a, radial point 122 lies at the tip of blade 106. Each blade 106 has a connecting link 130 extending from second end 110.

Connecting link 130 ends in mounting means 134, typically a mounting flange adapted for attachment to a linear bearing block, for interfacing with an actuation device, shown generally at 138. Actuation device 138 is capable of simultaneously moving blades 106 to alter the size of aperture 118.

5 Actuation device 138 includes actuation plate 142 which is coaxial with reference circle 114. Actuation plate 142 has eight equi-spaced radial slots 146. More generally, for every blade there will be a corresponding radial slot on actuation plate 142. Each radial slot 146 overlaps a mounting means 134 for a linear bearing block at the end of a connecting link 130. Each blade 106 is engaged to actuation plate 142 via a
10 cam follower bearing 150 disposed in radial slot 146 and attached to mounting means in slotted end 134.

Each bearing 150 extends from a linear slide 154. Linear slide 154 is mounted on a non-rotating plate 156 (shown in Figure 8). Linear slide 154 is constructed and arranged to slide along a line 158 which is parallel to the radius 126 on
15 which radial point 122 of blade 106 lies.

For the purposes of this disclosure, the term 'cam follower bearing' includes cam follower bearings, low friction rollers, roller bearings, needle roller bearings and a slipper block pivot mounted on a bearing and stub shaft. Figure 4b is a partial front view of an embodiment in which a slipper block is used. A side view of the
20 embodiment of Figure 4b taken along lines 4c-4c is shown in Figure 4c. Slipper block 150 resides in slot 146 of actuation plate 142. Slipper block 150 is mounted on stub shaft 151 which extends from connecting link 130. Desirably, bearings 153 will be present between shaft 151 and slipper block 150. Connecting link 130, in turn, is fastened to linear bearing block 212 via fasteners 214. Bearing block 212 is linearly
25 mounted on linear slide which is mounted on fixed plate 156. Linear motion is facilitated by the presence of bearings 216.

Cam follower bearing 150 may be replaced by any other suitable connecting member which can connect the slide and the link.

In use, as actuation plate 142 is rotated in a clockwise direction, the
30 clockwise motion of the actuation plate is translated into linear motion of each of linear slide 154 and blade 106 via bearing 150. Each blade 106 moves outward in a direction parallel to the radius 126 on which the radial point 122 of the blade 106 lies, resulting in the opening of aperture 118. As actuation plate 142 is rotated in a counterclockwise

direction, each blade 106 moves inward in a direction parallel to the radius 126 on which the radial point 122 of the blade 106 lies, resulting in the closing of aperture 118. As aperture 118 closes, a radially inward force is applied to a medical device disposed in the aperture. The actuation plate is rotated until the desired size reduction of the aperture and medical device has been achieved. Following the reduction, the actuation plate is rotated in the opposite direction to allow for removal of the medical device from the aperture.

The apparatus may be used to reduce the diameter of a suitable medical device such as those disclosed above or may be used to crimp a medical device to a catheter.

Another embodiment of the invention is shown in Figure 5a. Each blade 106, as shown in Figure 5a, has a connecting link 130 extending therefrom. Connecting link 130 is rigidly attached to blade 106. Connecting link 130 ends in an angled end 134 for interfacing with an actuation device, shown generally at 138. Actuation device 138 is capable of simultaneously moving blades 106 to alter the size of aperture 118.

Actuation device 138 includes a rotatable actuation plate 142 which is co-axial with reference circle 114. Rotatable actuation plate includes cam slots 146 which are not concentric with the axis of rotation, arcing inward. Each connecting link 130 is engaged to actuation plate 146 via a cam follower bearing 150 disposed in slot 146 and attached to both angled end 134 of connecting link 130 and to a linear slide 154. Linear slide 154 is mounted on a non-rotating plate similar to that shown in Figure 8. Linear slide 154 is constructed and arranged to slide along a radial line 158 on which radial point 122 of blade 106 lies.

Connecting link 130 may be bonded adhesively, welded, joined with a fastener or otherwise joined to blade 106. As shown in Figure 5a, a single screw 131 is used to connect link 130 to blade 106. Figure 5b shows a connecting link 130 including a right angle portion which is fastened to a blade 106 using two screws 131. Connecting link 130 and blade 106 may optionally be formed of a single piece of material.

Regardless of how the connecting member is joined to the blade, no movement of the blade relative to the connecting link is permitted.

In use, as actuation plate 142 is rotated in a clockwise direction, the clockwise motion of the actuation plate is translated into a linear outward motion of each of linear slides 154 and blades 106 via bearings 150 resulting in the opening of

aperture 118. The outward motion results from the radially outward arcing of cam slot 146. As actuation plate 142 is rotated in a counterclockwise direction, each blade 106, because of the radially inward arc of cam slots 146, moves inward in a direction parallel to the radius 126 on which the radial point 122 of the blade 106 lies, resulting in the
5 closing of aperture 118. As discussed above, as the aperture is decreased in size, a radial inward force is brought to bear against a medical device disposed in the aperture, thereby reducing the size of the medical device.

The embodiment of Figure 5a differs from the embodiment of Figure 4a in that in the embodiment of Figure 5a, the slide moves along the radial line on which
10 the radial point of the attached blade lies whereas in Figure 4a the slide moves parallel to the radial line. In both of the embodiments, each of the blades is constrained with two degrees of freedom to satisfy the condition that the movement of the tip be radial in accordance with the invention.

In the embodiments of Figures 4a and 5a, the slots in the actuation plate
15 are constructed and arranged to allow for a sufficient reduction in size of the aperture so that a medical device can be reduced in size to a desired diameter. Those of ordinary skill in the art will recognize other suitable actuation devices that may be used in the practice of this invention.

Desirably, in the above embodiments, the blades will be as long as or
20 longer than the medical device disposed within so that the medical device is uniformly reduced in size along its entire length.

This is illustrated in the embodiment of Figures 6 and 7 and further in Figures 3a and 3b in which blades 106 are disposed about a reference tube 160 to form a tubular aperture 162 whose size may be varied. Reference circle 114 is seen to lie along
25 reference tube 160. Each blade 106 is in communication with an actuation device such as that shown in Figures 4 or 5. The actuation device is capable of moving blades 106 to alter the size of tubular aperture 162. Each blade 106 includes a single line 166 which a) lies on a radial plane 170 of the reference tube 160 prior to movement of blade 106, and b) may be moved only along a radial plane 170 of reference tube 160 on
30 movement of blade 106. Desirably, reference tube 160 is cylindrical and exceeds the length of the medical device to be reduced in size.

Another embodiment of the invention is illustrated in Figures 8a and 8b. In the embodiment of Figures 8a and 8b, two non-rotating plates 156 are present, one at

each end of the apparatus. Each blade 106 is connected at first end 174 to a linear slide 154a via a connecting link 130a and at second end 178 to a linear slide 154b via a connecting link 130b. Linear slide 154a is mounted on non-rotating plate 156a and linear slide 154b is mounted on non-rotating plate 156b. The presence of the second
5 non-rotating plate 156b, linear slide 154b and connecting link 130b is optional but contributes to providing a rigid frame upon which the connecting links and associated blades may slide without misalignment relative to the reference circle.

Figures 8a and 8b illustrate the use of the inventive apparatus in various stages of the size reduction process. In Figure 8a, stent 180 has been placed in tubular
10 aperture 162 which is characterized by a diameter d_1 . In Figure 8b, the device has been actuated by rotating actuation plate 142 so as to move blades 106 inward. Aperture 162, as shown in Figure 8b is characterized by a diameter d_2 which is reduced relative to diameter d_1 . Stent 180 is seen to be of reduced diameter relative to its previous diameter as shown in Figure 8a.

15 Figure 8c differs from Figure 8b, only in that stent 180 has been crimped onto catheter 184 in Figure 8c.

Blades 106 may be made of any suitable, hard material including hardened steel. Desirably, the blades will be made of a material such as zirconia ceramic. Blades made of zirconia ceramic may be used without lubrication.
20 Furthermore, because of their low thermal conductivity, they may be used to create a highly insulated chamber suitable for cryogenic processing of martensite in nitinol stents.

Such an embodiment is shown in Figure 9. Stent 180 is disposed between blades 106 which can move inward in the direction of the arrows. Blades 106
25 are cooled by a first source of cooling fluid 184 located at first end 174 of blades 106. Although not shown, a second source of cooling fluid may be provided at second end 178 of blades 106 as well. The cooling fluid may be a liquid cryogenic. Exemplary cryogenics include liquid nitrogen, argon or carbon dioxide although other cryogenics may also be used. The cooling fluid may also be a chilled gas such as air. The cooling
30 fluid may also be a cooled inert gas such as nitrogen, argon or other inert gasses.

The aperture formed by the blades is a highly insulated chamber which is suitable for cryogenic processing of martensite in nitinol stents. The chamber is maintained at -80°C and a nitinol stent inserted therein. Upon equilibration of the

temperature of the stent, the blades are moved inward to reduce the diameter of the stent. The stent is thus reduced in diameter while being maintained in the martensitic state.

The embodiment of Figure 9 further has a loading plunger 188 for
5 loading a stent or other suitable medical device into the aperture. A sheath housing 192 which houses sheath 196 is provided at second end 178 of blades 106. Plunger 188 may be further used to transfer the stent after it has been reduced in diameter or size to sheath 196. Desirably, sheath 196 will have a slightly larger diameter than stent 180 following reduction in size of the stent. More desirably, the fit of the stent within the sheath will
10 be within about 1/32" and even more desirably, within about 1/64".

Where lengthy stents or other medical devices are to be reduced in size, the invention contemplates using one of the above described apparatuses with long blades to accommodate the stent. As an alternative, the invention also contemplates disposing two or more of such apparatuses sequentially to form one long aperture. The
15 two or more apertures may then be reduced in size simultaneously or consecutively. The arrangement of Figure 10 shows an embodiment with three devices 100a-c arranged sequentially. A stepped reduction in size may be achieved by placing a stent 180 or similar medical device in the apparatus and independently reducing each aperture 118 a-c to a desired size. To that end, the invention may provide particular utility in
20 manipulating bifurcated stents or other stents whose diameter varies along its length. The embodiment of Figure 10 shows the end portions of the stent being reduced in size prior to the middle portion of the stent. The device may also be operated so as to reduce the middle portion in size prior to the end portions or in any other sequence.

The invention contemplates yet another approach to reducing the
25 diameter of lengthy stents or similar medical devices, namely walking the stent through the apparatus. This may be accomplished by either moving the stent relative to the apparatus or moving the apparatus relative to the stent as shown schematically in Figure 11. To that end, stent 180 is inserted in device 100. Aperture 118a is reduced in size with blades 106a in turn reducing portion 180a of stent 180 in size. Aperture 118a is
30 then opened and aperture 118b reduced in size thereby reducing portion 180b of stent 180. Simultaneously, or shortly thereafter, sheath 196 is pushed by plunger 188 over the portion of the stent that has been reduced in size. Aperture 118b is opened and the stent

advanced in the apparatus. The process is repeated until the entire length, or the desired portion of the stent or medical device is reduced in size.

The reduction in size of the stent or other medical device may occur as part of a precrimping step or it may occur as part of crimping a stent onto a catheter and
5 desirably, onto a balloon disposed about a catheter. In a general sense, it may be used for manipulating a medical device and specifically, for applying a radial inward force to a medical device.

In another embodiment, the invention is directed to a method of manipulating a medical device. As part of the method, a medical device such as those
10 disclosed above is provided. The device has at least three blades capable of applying a radial inward force. The blades are disposed about a reference circle to form a shrinkable aperture. The blades are constructed and arranged such that each blade has only a single point which a) lies on the circumference of the reference circle prior to movement of the blade, and b) is moved along a radius of the reference circle on
15 movement of the blade. The medical device is placed into the shrinkable aperture and the blades simultaneously moved inward to apply a radial inward force to the medical device and thereby reduce the medical device in size, and desirably, in diameter. Following reduction in size of the medical device, the blades are simultaneously moved outward and the medical device removed from the aperture.

20 The inventive apparatus may also be incorporated into a blow molding tool to provide a variable size balloon mold as shown generally at 100 in Figure 12. The various parts of the apparatus of Figure 12 have been discussed in conjunction with Figures 8a-c and, with exception of balloon 181 and mold cavity ends 193, the reference numerals used in Figure 12 correspond to those used for Figures 8a-c. Mold cavity ends
25 193 may be provided in a variety of sizes and lengths to contain the balloon at each end. Desirably, the end molds will be adjustably mounted to a portion of the apparatus such as fixed plates 156 to provide for an adjustable length balloon mold.

The invention is also directed to a method for molding a medical balloon using the inventive apparatus described above. A balloon preform prepared through any
30 standard method is provided. The inventive mold, shown generally at 100 is also provided. Balloon 181 is inserted into aperture 162. Aperture 162 is optionally reduced to a predetermined size and the preform expanded using standard techniques. An

inflation fluid, for example, may be supplied to the preform and the preform expanded and heated. The balloon in its expanded state is shown in Figures 12.

More generally, the invention may be practiced by providing at least three movable blades disposed about a reference tube to form a shrinkable tubular aperture. The blades are constructed and arranged such that each blade has a single line which a) lies on the surface of the reference tube prior to movement of the blade, and b) is moved along a radial plane of the reference tube on movement of the blade. A balloon preform is placed into the shrinkable aperture. The aperture may be set at a predetermined size prior to or following insertion of the balloon therein. An inflation fluid is provided and the balloon preform inflated so that the preform expands to the size of the aperture. The preform may be heated during this inflation/blowing step. The inflation fluid is then removed from the thus formed balloon and the balloon removed from the apparatus.

The balloon may be also be molded in accordance with the method described in US 5,163,989, or in accordance with other methods as are known to those of ordinary skill in the art, substituting the instant apparatus for the standard balloon mold. Other patents which discuss balloon molding include US 5,807,520. Other references illustrating the materials and methods of making catheter balloons include: US 4,413,989 and US 4,456,000 to Schjeldahl et al, US 4,490,421, US Re 32,983 and Re 33,561 to Levy, and US 4,906,244, US 5,108,415 and US 5,156,612 to Pinchuck et al.

The use of the inventive apparatus as a mold allows for the blowing of a balloon to a predetermined size using a single adjustable size balloon mold thereby eliminating the need to have multiple molds of different sizes.

The invention further contemplates molding a balloon to a desired shape using a plurality of the inventive devices arranged sequentially. As an example of this, shown in Figure 13, a stepped balloon 181 may be prepared by arranging several devices 100a, 100b and 100c sequentially. A balloon preform is inserted in the aperture formed by the device. The aperture of each device may be preset at a desired size or may be reduced in size to a predetermined size after the balloon preform is inserted therein. The balloon may then be blow molded in accordance with any suitable blow molding technique known in the art.

The invention is also understood to be directed to embodiments employing various combinations of the features disclosed herein.

The above disclosure is intended to be illustrative and not exhaustive.

This description will suggest many variations and alternatives to one of ordinary skill in
5 this art. All these alternatives and variations are intended to be included within the scope of the attached claims. Those familiar with the art may recognize other equivalents to the specific embodiments described herein which equivalents are also intended to be encompassed by the claims attached hereto.

U.S. Application No. 09/401,218, from which this application claims
10 priority, is incorporated in its entirety herein by reference.

CLAIMS:

What is claimed is:

1. An apparatus for reducing in size a medical device comprising:
at least three coupled movable blades disposed about a reference circle to
5 form an aperture whose size may be varied, the aperture capable of being sized to
contain the medical device,
each blade in communication with an actuation device which is capable
of moving the blade to alter the size of the aperture,
wherein each blade includes a single radial point which
10 a) lies on a radial line of the reference circle prior to movement of
the blade, and
b) may be moved only along the radial line of the reference circle
on movement of the blade.
2. The apparatus of claim 1 wherein the blades are constructed and arranged to
15 form a shrinkable polygonal aperture.
3. The apparatus of claim 1 wherein the blades are constructed and arranged to
form a substantially circular aperture.
4. The apparatus of claim 1 wherein the actuation device comprises
a cam, and
20 a plurality of linear slide devices,
each of the linear slide devices in mechanical communication
with the cam,
and each blade is mounted to a linear slide device,
each linear slide device movable along an axis parallel to the radius on which the
25 radial point of the blade lies.
5. The apparatus of claim 1 comprising from 4 to 16 blades.
6. The apparatus of claim 5 comprising 8 blades.
7. The apparatus of claim 1 wherein the blades are disposed about a reference tube
to form a tubular aperture whose size may be varied, the reference circle lying along the
30 reference tube,
each blade in communication with an actuation device which is capable
of moving the blade to alter the size of the tubular aperture,
each blade includes a single line which

- a) lies on the circumference of the reference tube prior to movement of the blade, and
- b) may be moved only along a radial plane of the reference tube on movement of the blade.

- 5 8. The apparatus of claim 7 wherein the reference tube is a cylinder.
9. The apparatus of claim 1 in combination with a medical device which may be reduced in size, the medical device selected from the group consisting of stents, grafts stent-grafts and vena cava filters.
10. The apparatus of claim 1 wherein each blade has an inner end and an outer end,
10 the inner end being beveled to form a tip.
11. The apparatus of claim 13 wherein the single radial point on each blade is at the tip of the blade
12. An apparatus for applying a radial inward force to a medical device comprising:
at least three coupled movable blades disposed about a reference circle to
15 form an aperture whose size may be varied, the aperture capable of being sized to contain the medical device,
each blade in communication with an actuation device which is capable of moving the blade to alter the size of the aperture,
wherein each blade includes a single radial point which
20 a) lies on the circumference of the reference circle prior to movement of the blade, and
b) may be moved only along a radius of the reference circle on movement of the blade.
13. The apparatus of claim 12 with a medical device selected from the group
25 consisting of stents, grafts, stent-grafts and vena cava filters disposed in the aperture.
14. The apparatus of claim 13 wherein the medical device is disposed about a catheter.
15. An apparatus for crimping a medical device comprising:
at least three coupled movable blades disposed about a reference circle to
30 form an aperture whose size may be varied, the aperture capable of being sized to contain the medical device,
each blade in communication with an actuation device which is capable of moving the blade to alter the size of the aperture,

wherein each blade includes a single radial point which

a) lies on the circumference of the reference circle prior to movement of the blade, and

5 b) may be moved only along a radius of the reference circle on movement of the blade.

16. A method of manipulating a medical device comprising the steps of:
providing a medical device;
providing at least three blades capable of applying a radial inward force, the blades disposed about a reference circle to form a shrinkable aperture;
10 placing the medical device into the shrinkable aperture;
simultaneously moving the blades inward to apply a radial inward force to the medical device;
wherein the blades are constructed and arranged such that each blade has a single point which

15 a) lies on the circumference of the reference circle prior to movement of the blade, and
b) is moved along a radius of the reference circle on movement of the blade.

17. The method of claim 16 wherein the medical device is selected from the group
20 consisting of stents, stent-grafts, grafts and vena cava filters.

18. The method of claim 17 wherein the medical device is reduced in size.

19. The method of claim 17 wherein the medical device is characterized by a diameter and the diameter is reduced in size.

20. The method of claim 17 wherein the medical device is disposed about a catheter.

25 21. The method of claim 17 wherein the medical device is disposed about a balloon which is disposed about a catheter.

22. The method of claim 20 wherein a sufficient radial inward force is applied to crimp the medical device to the catheter.

23. The method of claim 21 further comprising the steps of:
30 simultaneously moving the blades outward; and
removing the medical device from the aperture.

24. A method of molding a balloon comprising the steps of:
providing a balloon preform;

providing at least three movable blades disposed about a reference tube to form a shrinkable tubular aperture, wherein the blades are constructed and arranged such that each blade has a single line which

- 5 a) lies on the surface of the reference tube prior to movement of the blade, and
- b) is moved along a radial plane of the reference tube on movement of the blade;

placing the balloon preform into the shrinkable aperture;

providing an inflation fluid; and

- 10 inflating the balloon preform so that the preform contacts the blades.

25. The method of claim 24 wherein the balloon is heated during the inflating step to form a balloon.

26. The method of claim 25 wherein the inflation fluid is removed from the formed balloon and the balloon is removed from the aperture.

1 / 15

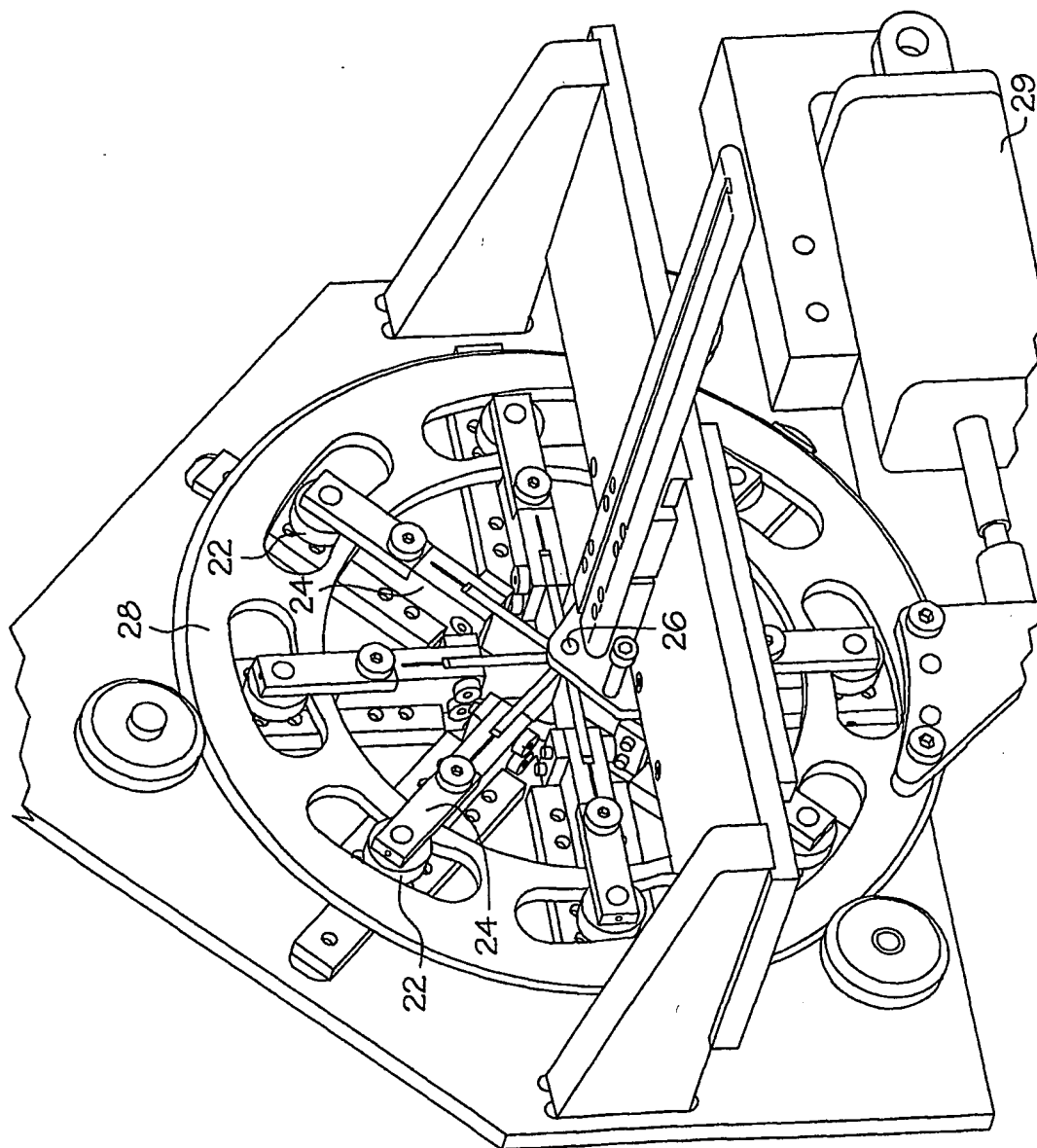


FIG. 1

2 / 15

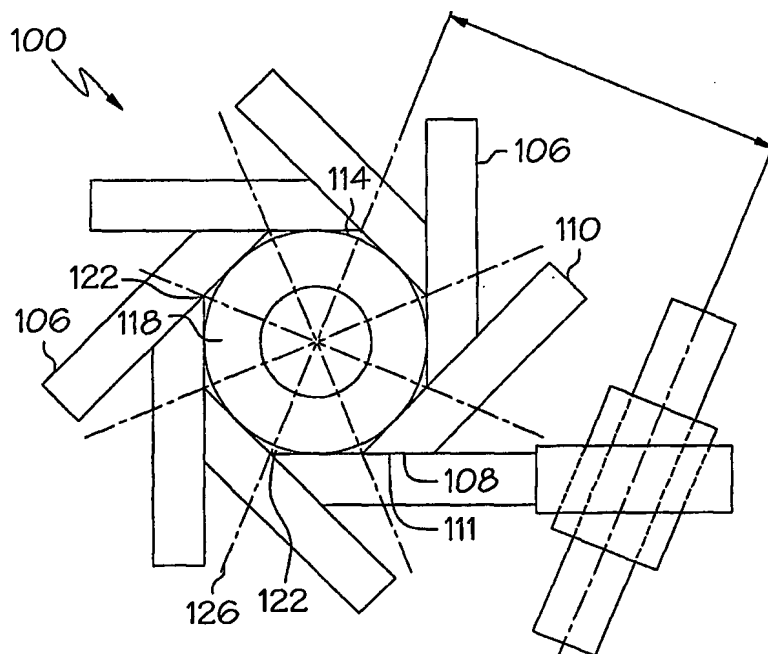


FIG. 2a

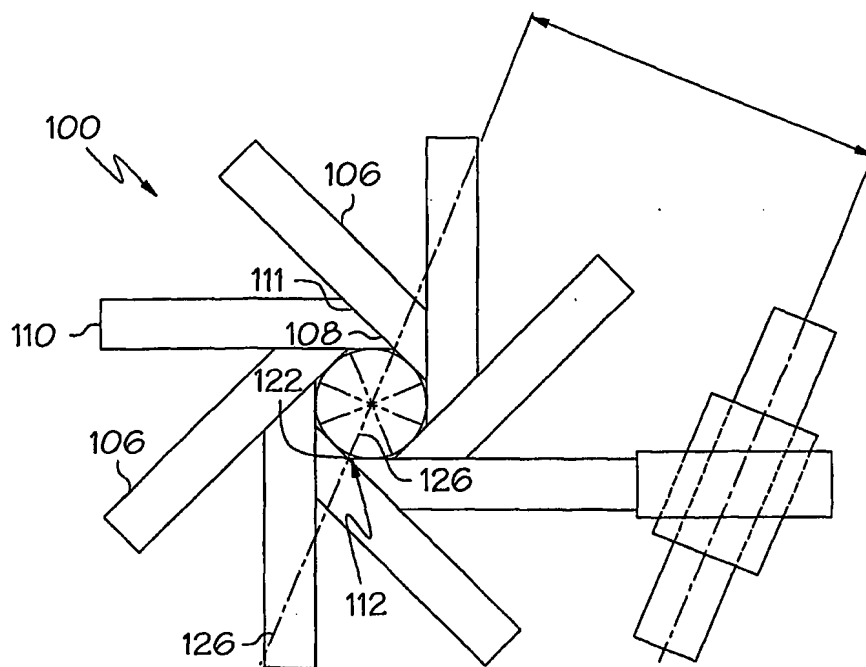


FIG. 2b

3 / 15

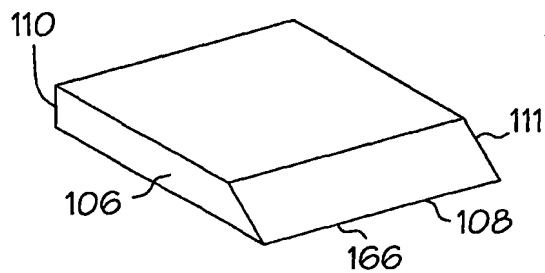


FIG. 3a

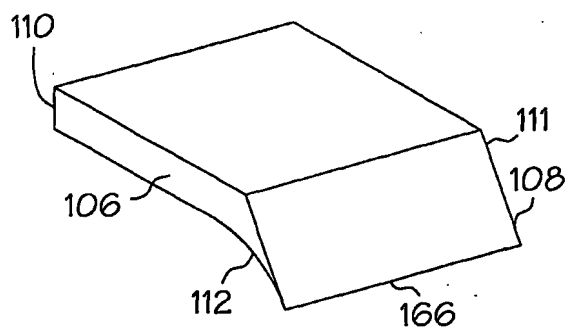


FIG. 3b

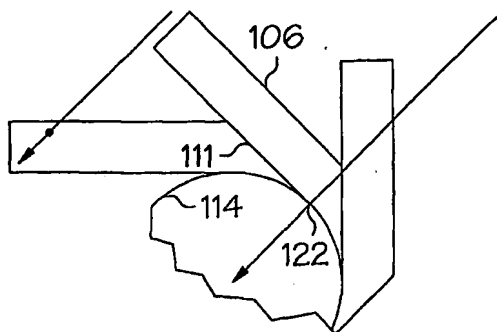


FIG. 3c

4 / 15

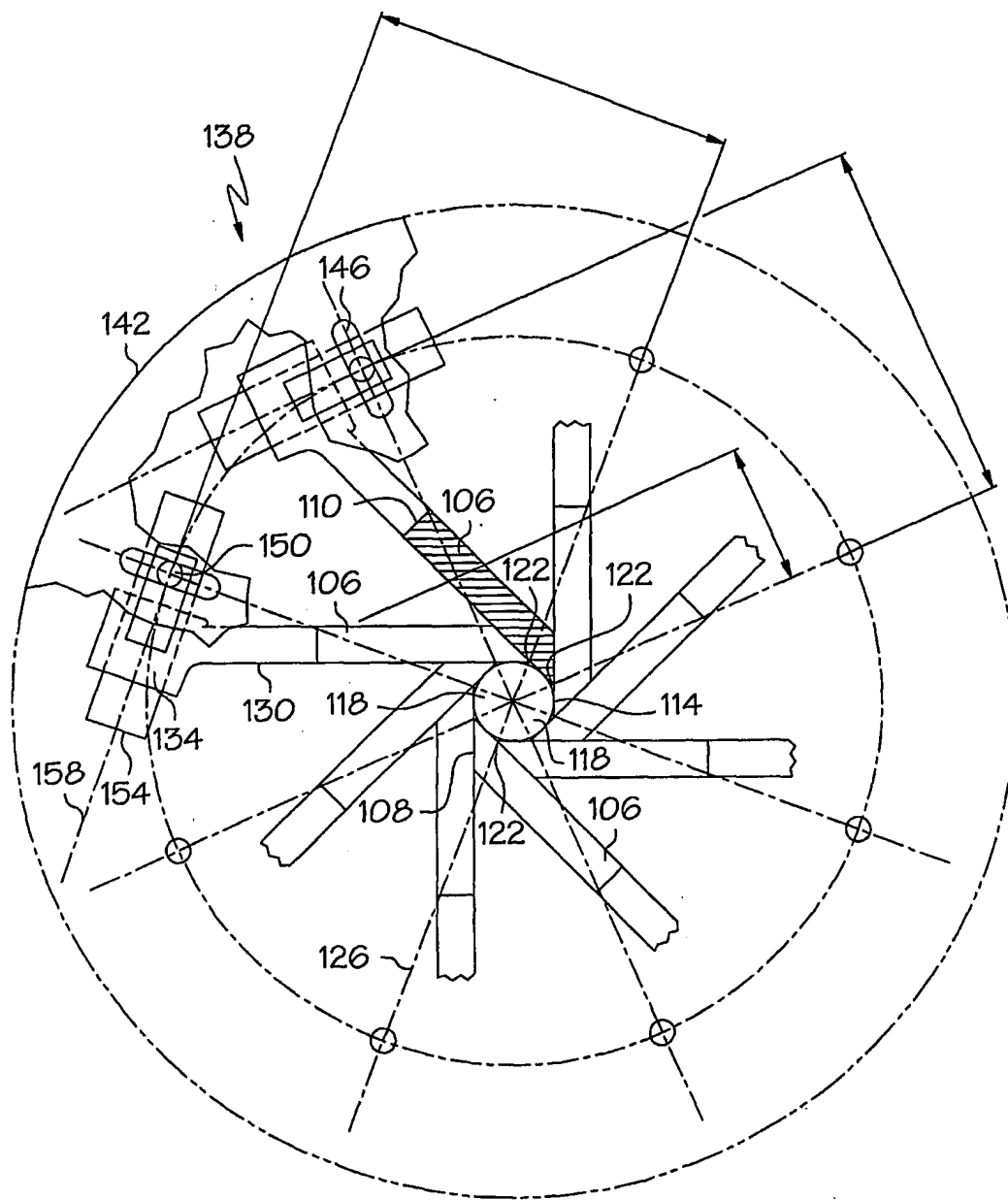


FIG. 4A

5 / 15

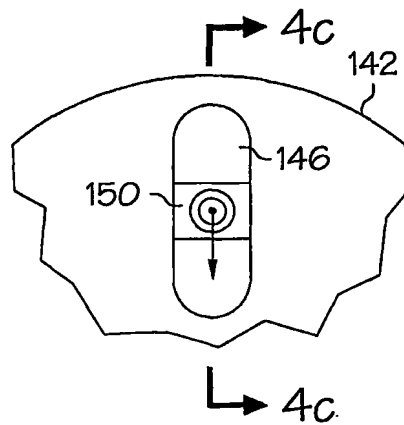


FIG. 4b

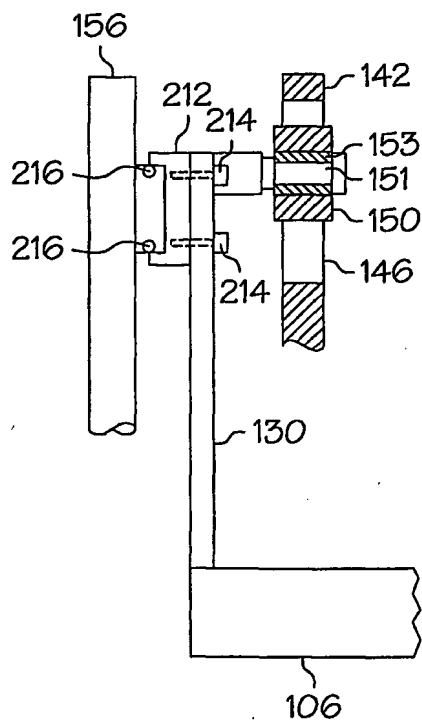


FIG. 4c

6 / 15

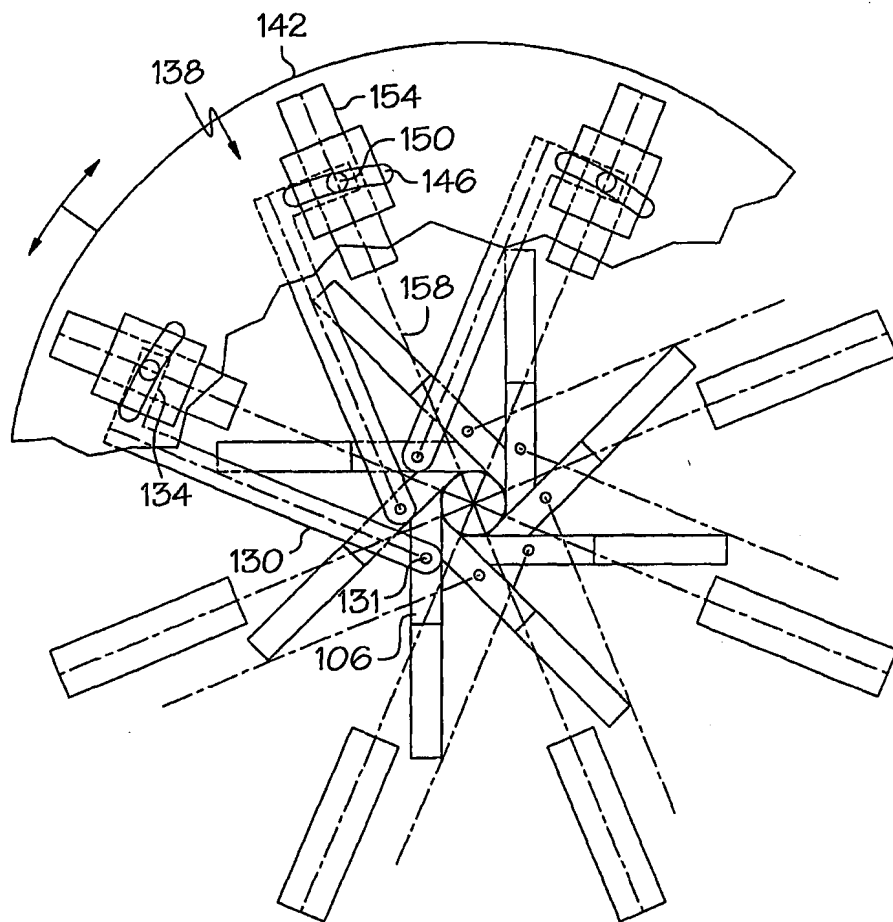


FIG. 5a

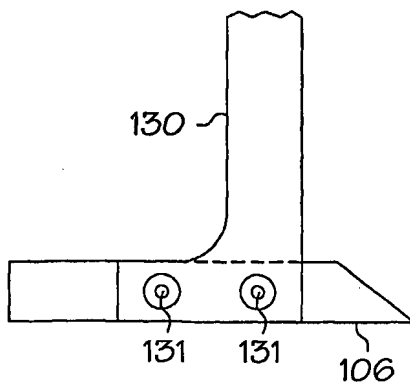


FIG. 5b

7 / 15

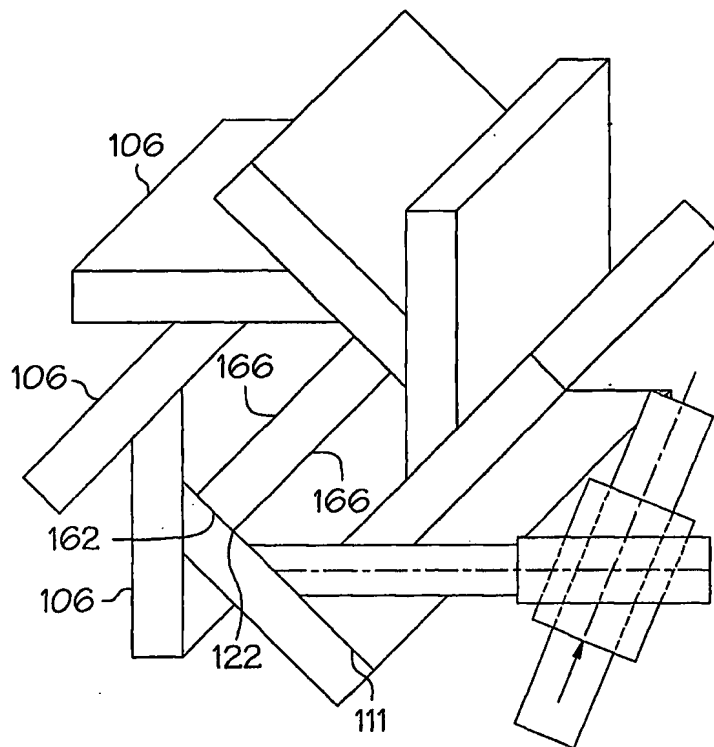


FIG. 6

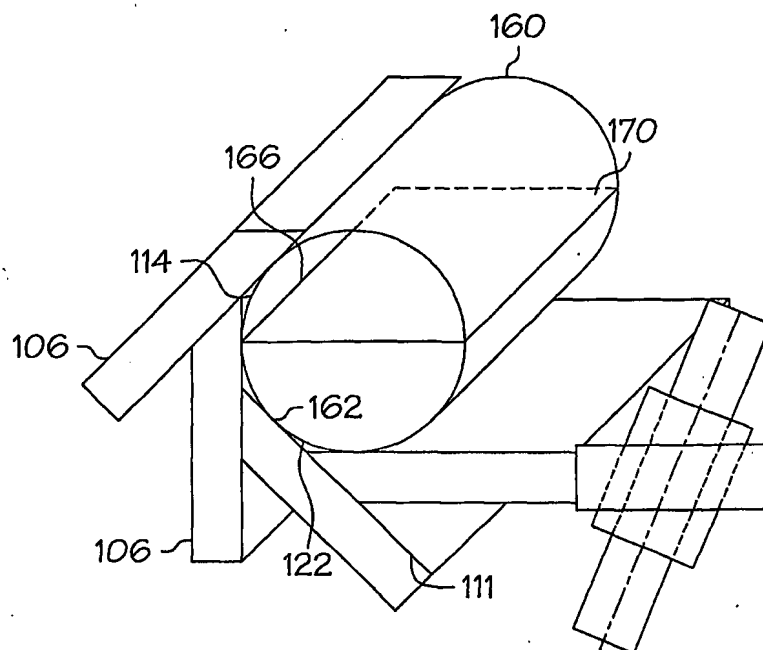


FIG. 7

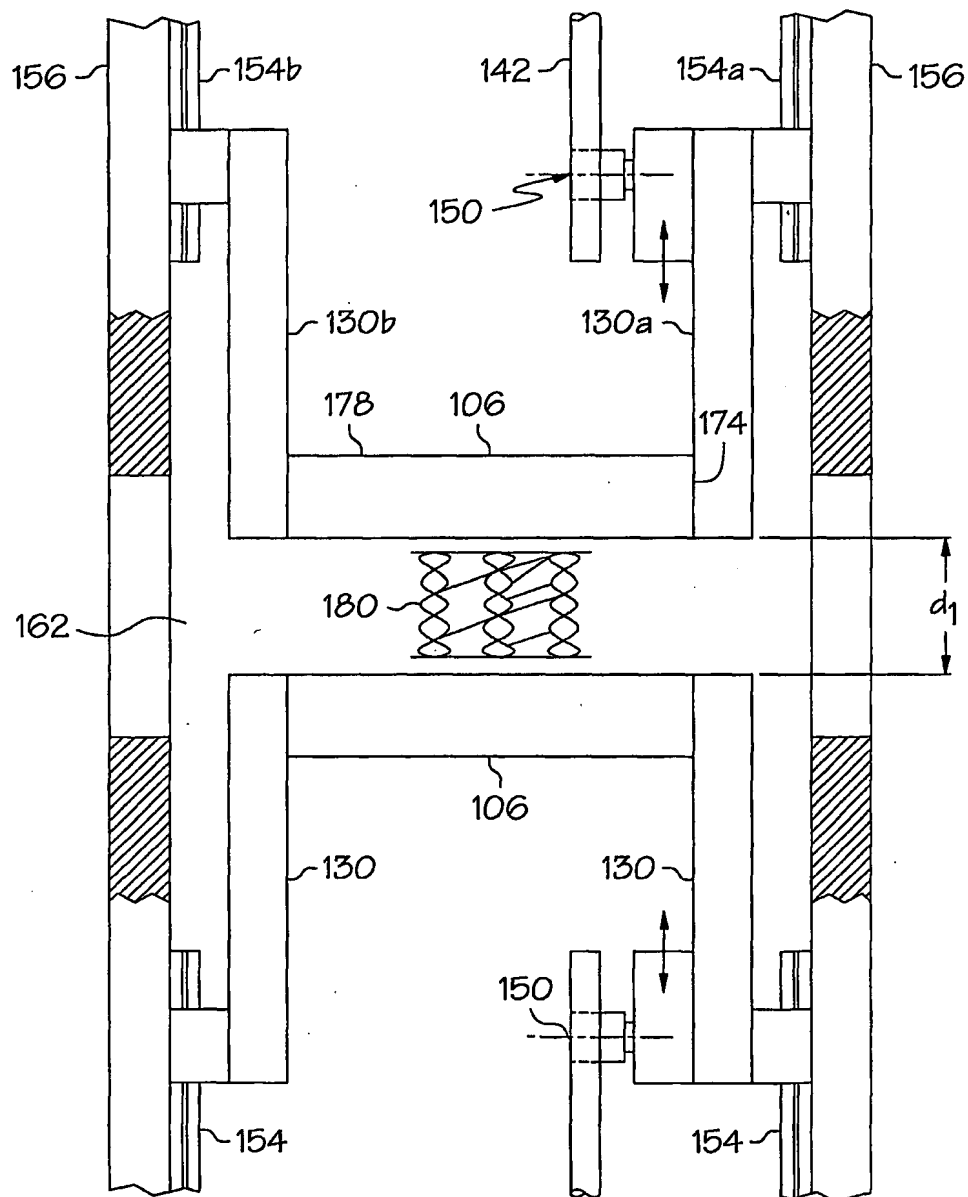


FIG. 8a

9 / 15

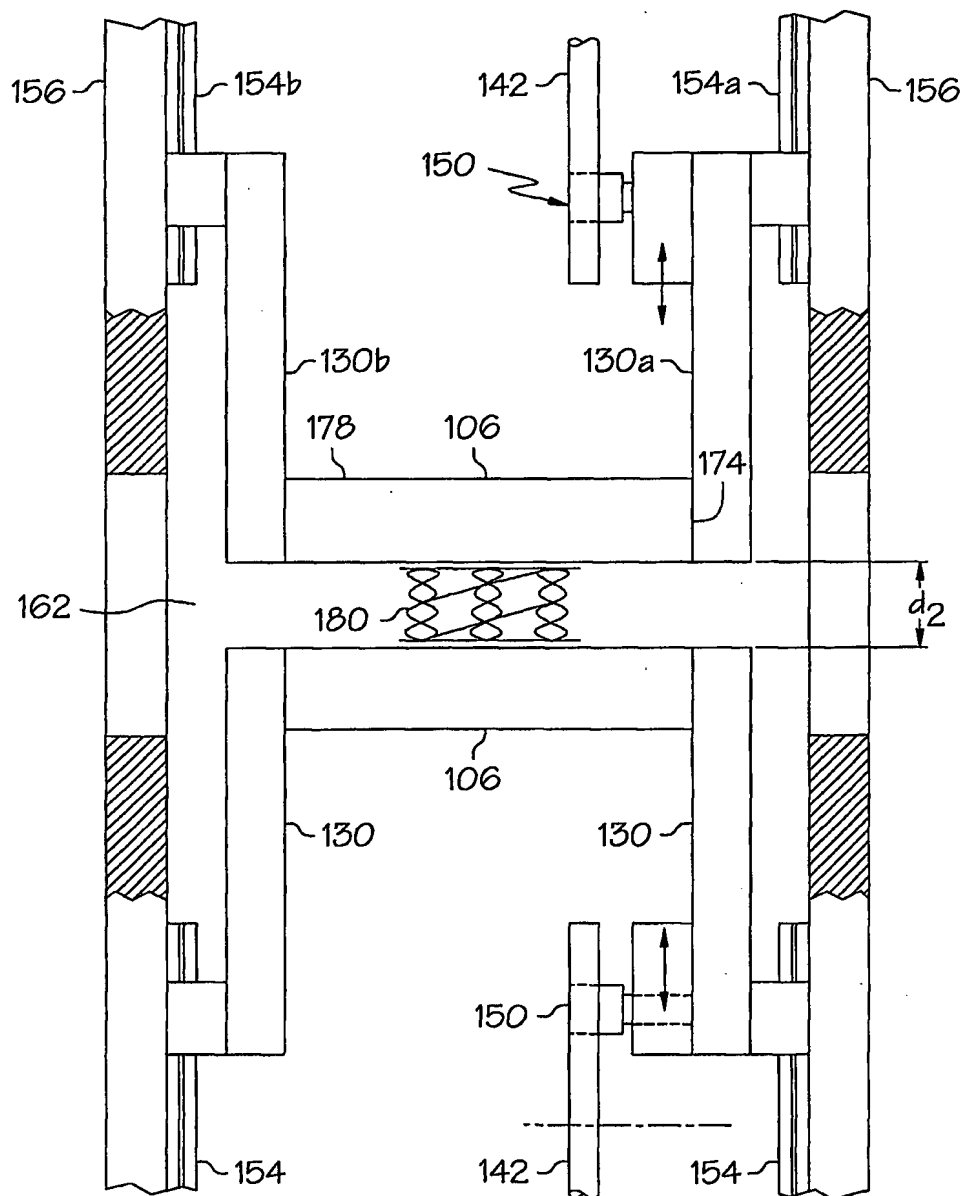


FIG. 8b

10 / 15

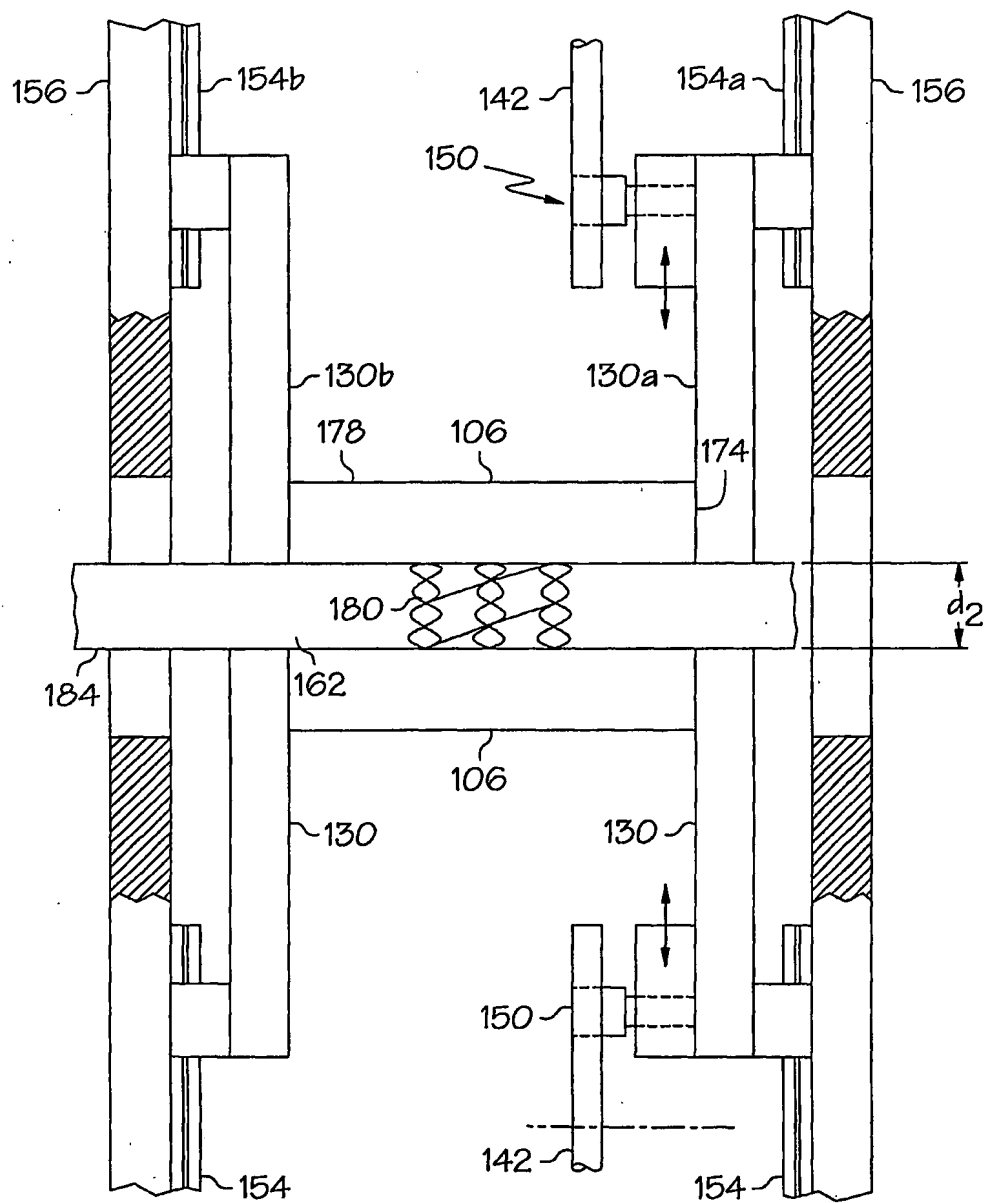


FIG. 8c

11 / 15

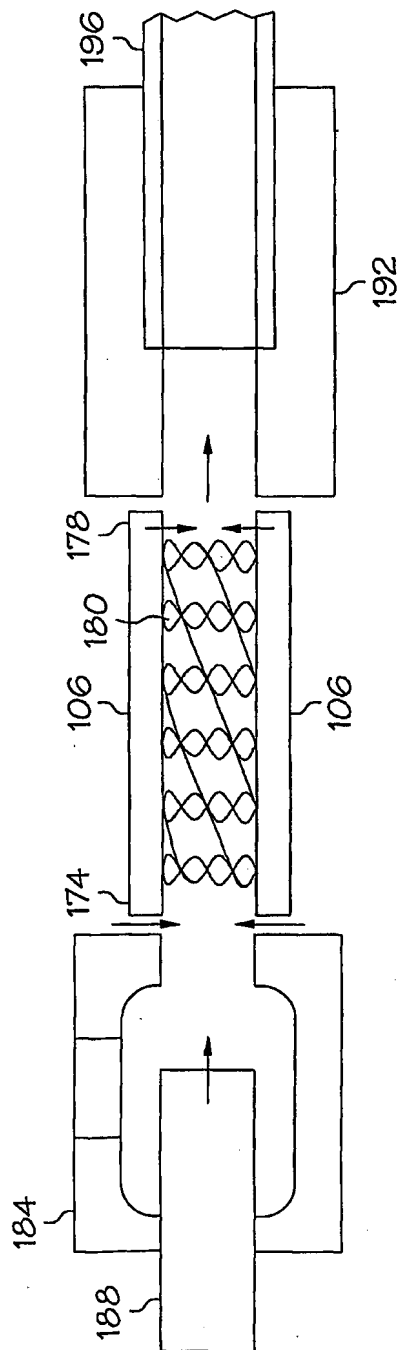


FIG. 9

12 / 15

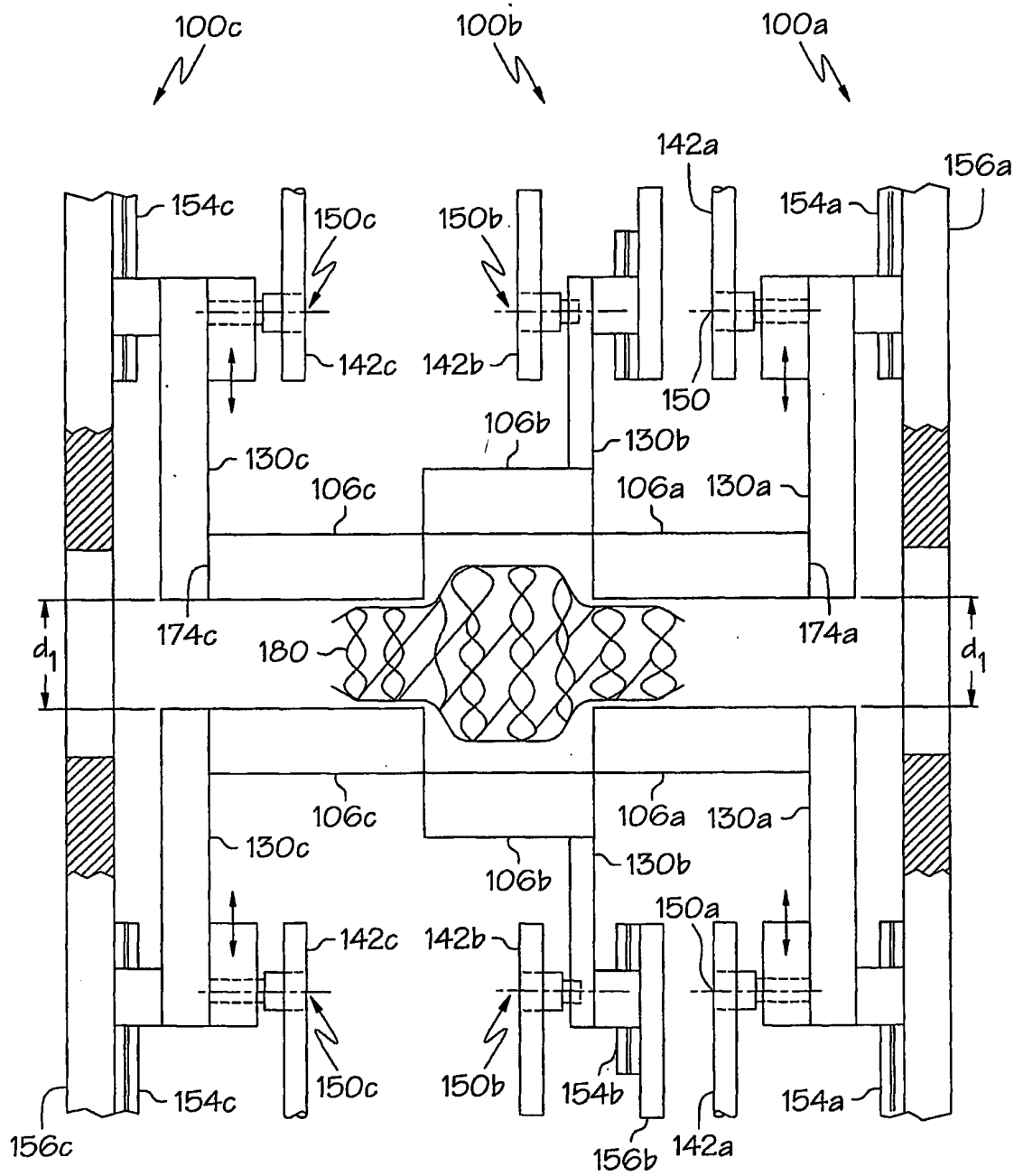


FIG. 10

13 / 15

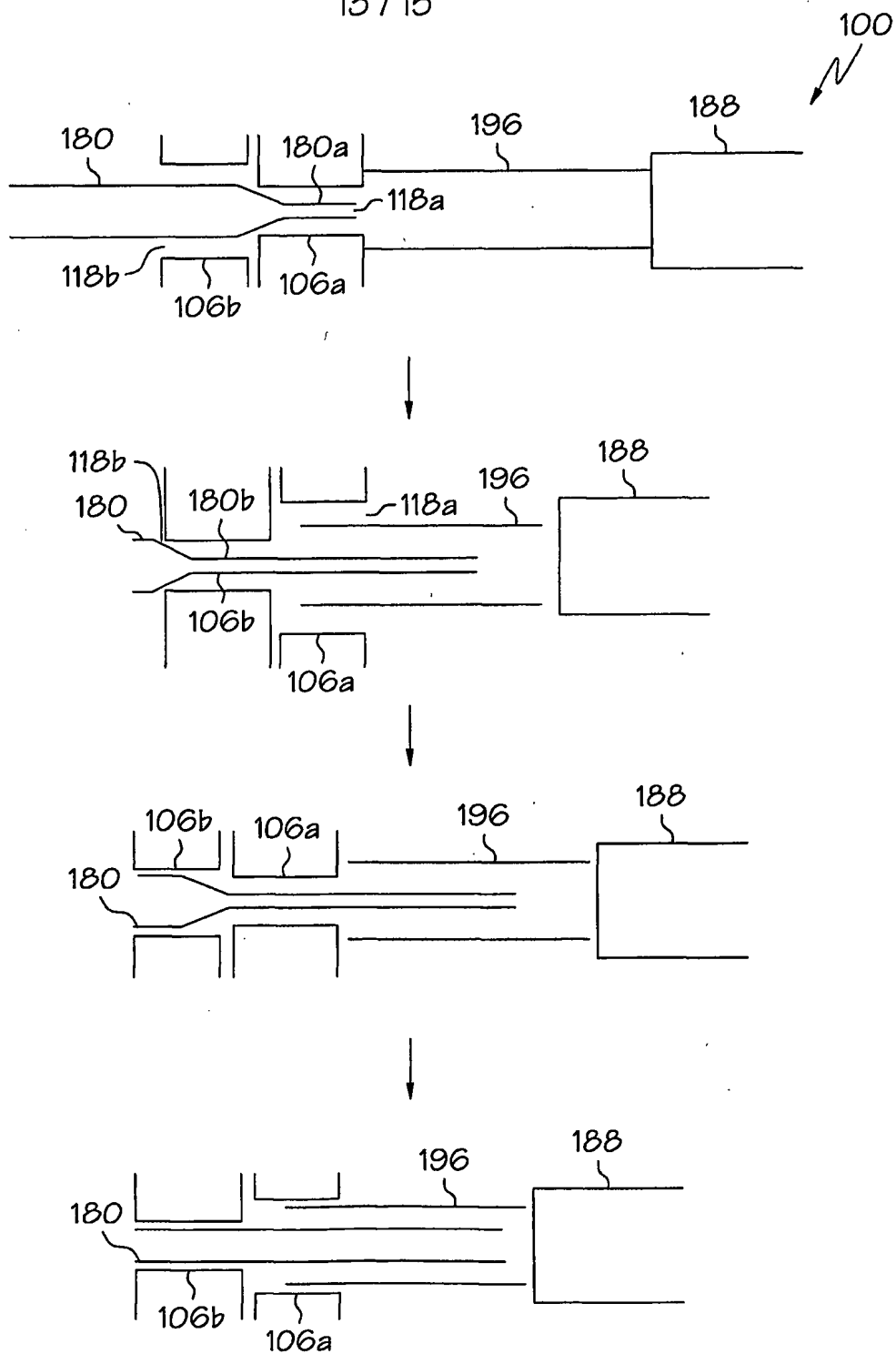


FIG. 11

14 / 15

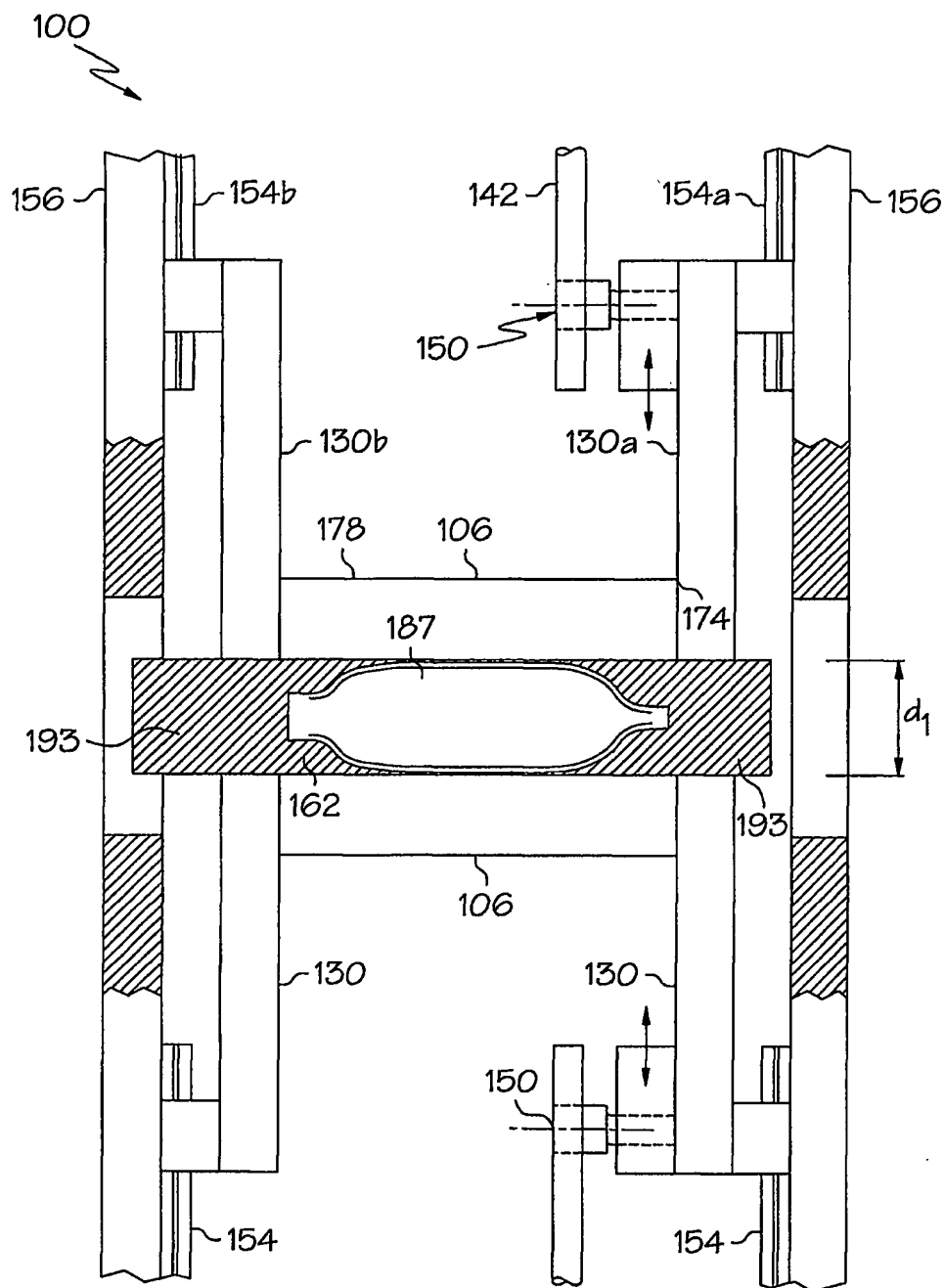


FIG. 12

15 / 15

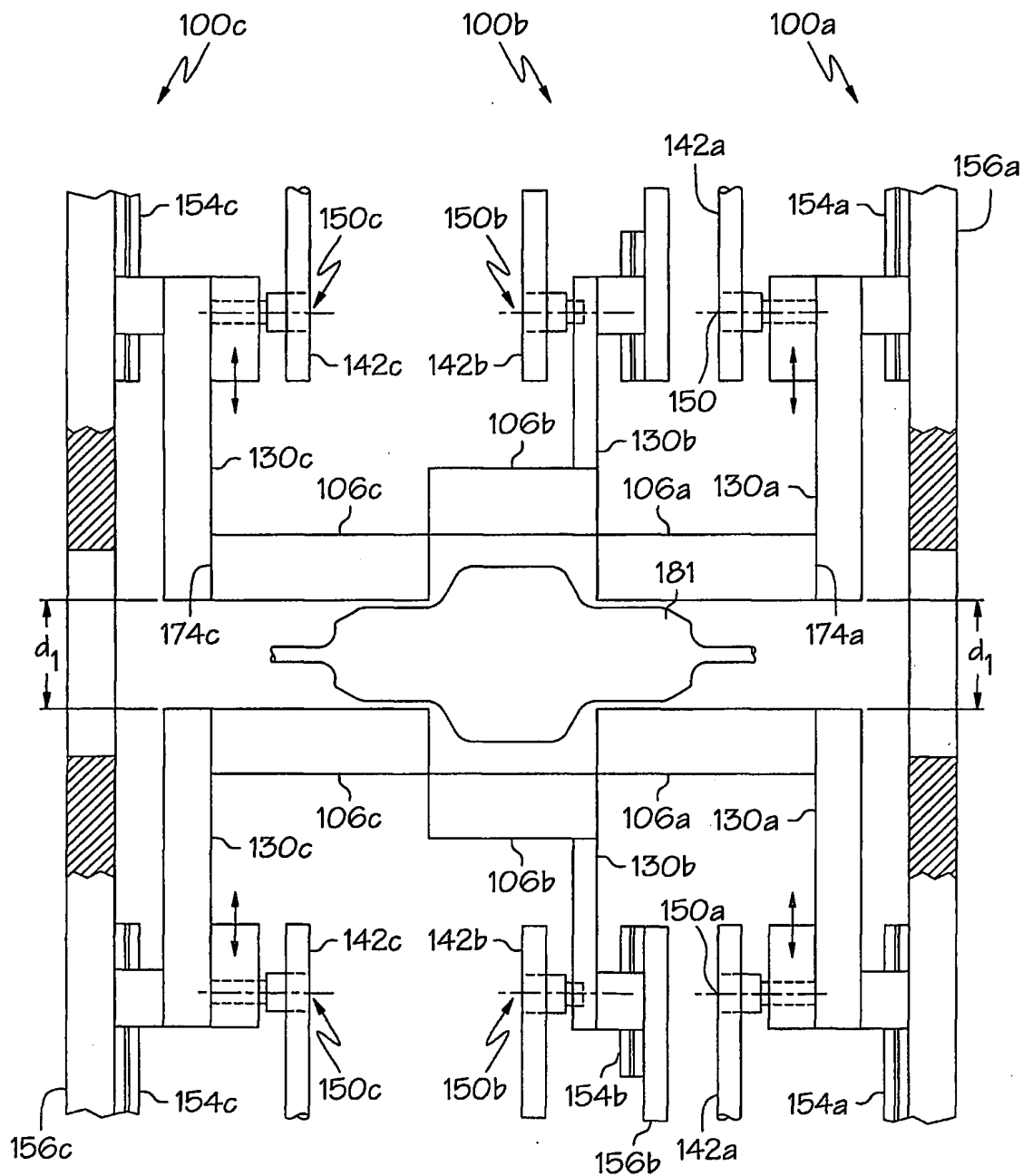


FIG. 13

INTERNATIONAL SEARCH REPORT

National Application No
PCT/US 00/25588A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61F2/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

WPI Data, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 935 952 A (ADVANCED CARDIOVASCULAR SYSTEMS, INC.) 18 August 1999 (1999-08-18) the whole document -----	1,12,15, 16

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

3 January 2001

Date of mailing of the international search report

10 April 2001 (10.04.01)

Name and mailing address of the ISA

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Authorized officer

Smith, C

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 00/25588

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-23

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-23

Apparatus and method for reducing in size a medical device.

2. Claims: 24-26

Method of moulding a balloon.

International Application No
PCT/US 00/25588

PCT/US 00/25588

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 935952 A	18-08-1999	US 6082990 A JP 11332996 A	04-07-2000 07-12-1999
